INFORMATION DISCLOSURE

STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number Filing Date First Named Inventor John Art Unit Examiner Name Unkn Attorney Docket Number		10826762		
		2004-04-16		
		Harper		
		2628		
		nown		
		P3352US1/119-0033US		

		Remove							
Examiner Initial*	r Cite No Patent Number		Kind Code ¹	Issue Date	Name of Patentee or Applicant of cited Document	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear			
	1	5877741		1999-03-02	Chee et al.				
	2	5877762		1999-03-02	Young et al.				
	3 6911984 2005-06-28 Sabella et al.		Sabella et al.						
If you wish to add additional U.S. Patent citation information please click the Add button.									
			U.S.P	ATENT APPLI	CATION PUBLICATIONS	Remove			
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publication Date	Name of Patentee or Applicant of cited Document	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear			
	1	20020067418	A1	2002-06-06	Hiroakı				
	2	20020093516	A1	2002-07-18	Brunner et al.				
	3	20040032409	A1	2004-02-19	Martin Girard				

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10826762		
Filing Date		2004-04-16		
First Named Inventor	John	John Harper		
Art Unit		2628		
Examiner Name	Unknown			
Attorney Docket Number		P3352US1/119-0033US		

	4	2005	0088452	A1	2005-0	4-28	Hanggie et al.						
If you wis	h to a	dd add	fitional U.S. Publis	shed Ap	plication	citatio	n information p	leas	se click the Add	butto	n. Add		
					FOREIG	GN PAT	ENT DOCUM	ENT	rs		Remove		
Examiner Initial*	Cite No	Fore	ign Document ber ³	Country Code ²		Kind Code ⁴	Publication Date	App	me of Patentee plicant of cited cument		where Re	or Relevant	т5
	1												
If you wish to add additional Foreign Patent Document citation information please click the Add button Add													
NON-PATENT LITERATURE DOCUMENTS Remove													
Examiner Cite initials* Cite (Dook, magazine, journal, serial, symposium, catalog, etc.), date, pages(s), volume-issue number(s), publisher, city and/off country where published.									Ţ5				
	1 International Search report dated March 8, 2006 (PCT/US 05/019108; 119-0032WO)								Z				
If you wish to add additional non-patent literature document citation information please click the Add button Add													
EXAMINER SIGNATURE													
Examiner	Signa	ture							Date Conside	red			

See Kins Codes of USPTO Patient Documents at year USPTO_CODE or MREP 901.04. ² Eatins office that insued the document, by the Nov-liter code (WIPO Standard ST.3). ² Standard ST.3). ² Standard ST.3 ² Standard ST.16 if possible. ³ Applicant is to place a check mark here if Proplet Imageage threatedors is attached.

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number			10826762		
Filing Date First Named Inventor John			2004-04-16		
		John	Harper		
	Art Unit		2628		
	Examiner Name	Unkn	own		
Attorney Docket Number			P3352US1/119-0033US		

CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s);

That each item of information contained in the information disclosure statement was first cited in any communication of forms a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(a)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.56(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.97(eV).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- .7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/Coe F. Miles/	Date (YYYY-MM-DD)	2006-05-09
Name/Print	Coe F. Miles	Registration Number	38559

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 3T CFR.

1.14. This collection is estimated to take I hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operatment of Commence, P. O. Box 1430, Alexandriu, V.S. 2213.1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.A. 2213.1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiation.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record perfains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designe, cuting an inspection of records concluded by GSAs a part of that apency's responsibility to recommend improvements in records management practices and programs, under suthority of 4d U.S.C. 2004 and 2006. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a pleath pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via stife of manaplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.